INFORMATION DISCLOSURE
STATEMENT BY APPLICANT
(Not for submission under 37 CFR 1.99)

Application Number		
Filing Date		
First Named Inventor	Alfon	sa CASAROTTO
Art Unit		
Examiner Name		
Attorney Docket Numb	er	06UVB004

					U.S.I	PATENTS			Remove	
Examiner Initial*	Cite No Patent Number		Kind Code ¹			Name of Pate of cited Docu	entee or Applicant iment	Pages,Columns,Lines where Relevant Passages or Relevant Figures Appear		
	1	5188119		1993-0	2-23	Sutherland		Col. 9, I. 67; Col. 10, I. 42; F		i. 1,9
	2	3326206 1967-06-20 Barr et al.		Col. 3, I. 16-63; Figs. 1-5						
	3	5938622		1999-0	8-17	Chen		All		
If you wis	h to a	dd additional U.S. Pater	t citatio	n inform	ation pl	ease click the	Add button.		Add	
			U.S.P	ATENT	APPLIC	CATION PUBL	LICATIONS		Remove	
Examiner Initial*	Cite No	Publication Number	Kind Code ¹	Publica Date	ition	Name of Pate of cited Docu	entee or Applicant iment	Releva	Columns,Lines where ant Passages or Relev s Appear	
	1									
If you wisl	h to a	dd additional U.S. Publi	shed Ap	plication	citation	n information p	olease click the Ad	d button	Add	
				FOREIG	SN PAT	ENT DOCUM	ENTS		Remove	
Examiner Initial*	Cite No	Foreign Document Number ³	Country Code ²		Kind Code4	Publication Date Name of Patente Applicant of cited Document		e or	Pages,Columns,Lines where Relevant Passages or Relevant Figures Appear	т
	1	0732077	EP		A	1996-09-18	von Froreich		Col. 6, L. 20, Col. 7, L. 28; Figs. 4, 5	

	Application Number		
NEODIA TION DIOOL COURT	Filing Date		
INFORMATION DISCLOSURE STATEMENT BY APPLICANT	First Named Inventor Alfon		sa CASAROTTO
(Not for submission under 37 CFR 1.99)	Art Unit		
,	Examiner Name		
	Attorney Docket Numb	or	OSI IVRODA

If you wisl	h to a	dd add	ditional Foreign Patent Document citation information please click the Add b	utton	Add		
			NON-PATENT LITERATURE DOCUMENTS		Remove		
Examiner Initials* Cite (locken name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, sorial, symposium, catalog, etc), date, pages(s), volume-issue number(s), publisher, city and/or country where published.							
	1						
If you wisi	h to a	dd add	ditional non-patent literature document citation information please click the	Add bu	tton Ad	d	-
			EXAMINER SIGNATURE				
Examiner	Signa	ture	Date Consider	ed			
			f reference considered, whether or not citation is in conformance with MPEP presence and not considered. Include copy of this form with next communics				

1 See Kin Codes of USPTO Patent Documents at wear USPTO_CODY or MPEP 90.16.2 Either office that issued the document, by the Involved (WIPO Standard ST3).3 "For Just planese petral for contents, the architecture of the Superior representations preceded he serial representations are the petral document. 4 Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST.16 if possible. 3 Applicant is to place a check mark here if Enginth Imagings the resistation is attached.

Application Number Fing Date Fing Date Fing Date Fing Market Parket Park

CERTIFICATION STATEMENT

Please see	37	CFR	1.97	and	1 98	to make	the	appropriate	selection(s)	r

That each item of information contained in the information disclosure statement was first cited in any communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of the information disclosure statement. See 3.7 CFR 197(eV1).

OR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing the certification after making reasonable inquity, no item of information contained in the information disclosure statement was known to any individual designated in 37 CFR 156(c) more than three months prior to the filing of the information disclosure statement. See 37 CFR 197(s)(c).

- See attached certification statement.
- Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.
- .7 None

SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

_			
Signature	/Franco A. Serafini/	Date (YYYY-MM-DD)	2006-05-03
Name/Print	Franço A. Serafni	Registration Number	52207

This collection of information is required by 37 CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is to file fand by the USPTO to process) an application. Confidentiality is governed by \$5 U.S. C.12 and 37 CFR 1.14. This collection is estimated to take 1 hour to complete, including gathering, preparing and submitting the completed application from the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patient and Tradenant's Office, u.S. Operatment of Commence, P. 0. Bot 1450, Alexandria, V.32.11.450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, V.32.211.41.60.

Privacy Act Statement

The Privacy Act of 1974 (P.L. 93-579) requires that you be given certain information in connection with your submission of the attached form related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, please be advised that (1) the general authority for the collection of this information is \$3 U.S.C. 2(b)(2); (2) famishing of the information solicided is voluntary, and (3) the principal purpose for which the information is used by the U.S. Patient and Trademan Kollie is to process and/or examine your submission related to a patient application or patient. If you do not furnish the requested process and/or examine your submission related to a patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested to the patient process and/or examine your submission, which may related that the patient process and/or examine your submission, which may

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these record s.
 - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement neodiations.
 - A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record partains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
 - A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974. as amended, pursuant to 5 U.S.C. 552a(m.).
 - A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
 may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
 to the Patent Cooperation Treaty.
 - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
 - 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designed, uturing an inspection of records conducted by GSA as part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 44 U.S.C. 2504 and 2506. Such disclosure shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make relevant for the state of the s
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the application pursuant to 35 U.S. C. 12(b) or issuance of a patient pursuant to 35 U.S. C. 15.1 Further, a record may be disclosed, subject to the limitations of 37 CFR.114, as a routine use, to the public if the record via flori of mapplication which became abandoned or in which the proceedings were terminated and which application is referenced by either a published application, an application open to public inseptions or an issued patient.
 - A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.